Exhibit C

IN THE UNITED STATES DISTRICT COURT FOR THE SOUTHERN DISTRICT OF NEW YORK

FEDERAL TRADE COMMISSION; STATE OF NEW YORK; STATE OF CALIFORNIA; STATE OF ILLINOIS; STATE OF NORTH CAROLINA; STATE OF OHIO; COMMONWEALTH OF PENNSYLVANIA; and COMMONWEALTH OF VIRGINIA,

Plaintiffs,

v.

VYERA PHARMACEUTICALS, LLC; PHOENIXUS AG; MARTIN SHKRELI, individually, as an owner and former director of Phoenixus AG and a former executive of Vyera Pharmaceuticals, LLC; and KEVIN MULLEADY, individually, as an owner and director of Phoenixus AG and a former executive of Vyera Pharmaceuticals, LLC,

Defendants.

Case No. 1:20-cv-00706-DLC

Plaintiffs' Disclosure of Opening Experts

Plaintiffs disclose the experts identified below who will provide opening expert reports on April 12, 2021.

1. C. Scott Hemphill, Ph.D., J.D.

Based on his experience and analysis of the relevant evidence, Professor Hemphill will opine on the relevant product market of FDA-approved pyrimethamine products, Defendants' monopoly power in the relevant market, the harm caused by Defendants' anticompetitive conduct, the lack of any procompetitive justification for Defendants' anticompetitive conduct, and the amount of Defendants' ill-gotten gains from their anticompetitive conduct. Professor Hemphill's curriculum vitae is attached as Exhibit A.

Professor Hemphill is available to provide deposition testimony on his expert opinions on July 8, July 15, or July 22.

2. W. David Hardy, M.D.

Based on his experience and analysis of the relevant evidence, Dr. Hardy will opine on toxoplasmosis and the serious health risks it poses, the historical trajectory and current state of toxoplasmosis in the United States, the diagnosis and treatment of toxoplasmosis, the unique role of Daraprim and other FDA-approved pyrimethamine products for the treatment of toxoplasmosis (including its status as the preferred therapy), and why trimethoprimsulfamethoxazole and compounded pyrimethamine products are not readily interchangeable with Daraprim for the treatment of toxoplasmosis. Dr. Hardy's curriculum vitae is attached as Exhibit B.

Dr. Hardy is available to provide deposition testimony on his expert opinions on July 27, July 29, or August 3.

3. James R. Bruno

Based on his experience and analysis of the relevant evidence, Mr. Bruno will opine on API sourcing and development, the CMC requirements for FDA approval of an ANDA, the availability of pyrimethamine API suppliers to supply the U.S. market in the relevant time period, the use of exclusive and backup API supply contracts, and Phoenixus's negotiations and contracts with Fukuzyu and RL Fine. Mr. Bruno's curriculum vitae is attached as Exhibit C.

Mr. Bruno is available to provide deposition testimony on his expert opinions on July 22, July 28, or July 29.

4. Edward V. Conroy

Based on his experience and analysis of the relevant evidence, Mr. Conroy will opine on: the distribution of pharmaceuticals and the role of specialty distribution; the historical distribution of Daraprim, including the use of specialty distribution for Daraprim; the sourcing of products in specialty distribution for use in bioequivalence testing; how Vyera's restrictive distribution system benefited the distributors of Daraprim financially; the role of data aggregators in the identification of drugs or generic drugs for development; and the uniqueness of Vyera's agreements and practices to prevent the sale of its distributors' Daraprim data. Mr. Conroy's curriculum vitae is attached as Exhibit D.

Mr. Conroy is available to provide deposition testimony on his expert opinions on July 14, July 21, or July 27.

Dated: February 12, 2021 Respectfully submitted,

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CERTIFICATE OF SERVICE

I hereby certify that on February 12, 2021, I served a true and correct copy of the foregoing **Plaintiffs' Disclosure of Opening Experts** on all counsel of record in this action via electronic mail.

Dated: February 12, 2021 Respectfully Submitted,

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